

Root Disease, Associated with *Verticicladiella alacris*, of Pines in South Africa

M. J. WINGFIELD, Plant Pathologist, Plant Protection Research Institute, and P. S. KNOX-DAVIES, Professor of Plant Pathology, University of Stellenbosch, Stellenbosch, 7600, South Africa

ABSTRACT

WINGFIELD, M. J., and P. S. KNOX-DAVIES. 1980. Root disease, associated with *Verticicladiella alacris*, of pines in South Africa. *Plant Disease* 64:569-571.

A disease of *Pinus pinaster* and *P. radiata* associated with root infection by *Verticicladiella alacris* was common at roadsides, and wounding appeared to be necessary for infection. All infected trees were colonized by the European bark beetle, *Orthotomicus erosus*, and *V. alacris* was isolated from *Hylastes* spp. in diseased roots.

Verticicladiella wagnerii Kendrick, *V. procera* Kendrick, and other *Verticicladiella* spp. are known root pathogens in other parts of the world (6,15,17,18). Infected trees are attractive to bark beetles (Coleoptera: Scolytidae) (1,9,11-13), which have been recorded as possible vectors of *V. wagnerii* (5). During a preliminary survey of diseases of exotic forest trees in South Africa, *Verticicladiella alacris* Wingfield & Marasas (19) was found associated with the roots of diseased *Pinus pinaster* Ait. and *P. radiata* D. Don. This paper describes the disease and its association with bark beetles.

MATERIALS AND METHODS

Diseased *P. pinaster* and *P. radiata* trees were examined in 13 centers in the following areas of the Western Cape Province: Cape Peninsula (Tokai State Forest), Franschoek (La Motte State Forest), and Grabouw (Grabouw and Lebanon State Forests).

Root systems were excavated and sampled. A sledge microtome was used to section infected roots, and isolations were made on half-strength malt extract agar (MEA) (10 g of Difco Malt extract and 15 g of Difco Bacto Agar per liter). Cultures were routinely maintained at 24 C.

One lateral root on each of 10 10-yr-old *P. pinaster* trees was inoculated by removing a 9-mm cambial disk with a cork borer and replacing it with a disk from a 2-wk-old culture of *V. alacris* on MEA. Inoculation points were covered with moist paper towels and plastic film, and the soil was replaced. Two-month-old *P. pinaster* and *P. radiata* seedlings (20 of each species) were inoculated by tying to their taproots, 20.0 × 3.0 mm wooden dowels, boiled in potato dextrose broth (200 g of potatoes and 15 g of dextrose per liter), and were colonized for 2 mo by *V. alacris*. Seedlings were repotted and kept in a glasshouse

maintained between 15 and 25 C. Equal numbers of control inoculations were made.

Naturally infected trees in all areas were examined for bark beetles. Beetles found in the roots and root crowns of trees in an infection center in the Grabouw State Forest were surface sterilized in a commercial sodium hypochlorite solution (1% available chlorine) with Tween 80 added, rinsed in sterile distilled water, and squashed onto MEA containing 4% vancomycin.

RESULTS

Symptoms. Infection centers in 10- to 15-yr-old plantings of *P. pinaster* and *P. radiata* ranged from a few trees to 2 ha in size. Many occurred at roadsides (Fig. 1). Trees showed reduced terminal growth and wilting, with chlorosis and death of the needles, which were retained after death. According to foresters in the area, trees took 2-3 yr to die, and during this period an abnormal number of cones was produced (Fig. 1). Infected roots and root crowns showed heavy impregnation of resin and dark blue discoloration with staining in areas parallel to the annual rings (Fig. 2). Staining was never seen above the root crown area. Symptoms on both *Pinus* spp. were similar, except that *P. radiata* roots stained darker and exuded more resin. Microtome sections through infected wood showed dark hyphae of *V. alacris* in the tracheids.

Natural regeneration (*P. pinaster*) replaced dead trees in infection centers, but many of the young trees were also diseased. In these trees healthy roots in contact with diseased roots developed lesions at the point of contact, and resin exuded from the bark of the roots (Fig. 3), which also stained dark blue (Fig. 4).

Inoculation tests. All roots inoculated after wounding exuded resin, and dark lesions similar to those observed in naturally infected trees developed in the sapwood. Six months after inoculation, lesions had extended an average of 10.0 cm on either side of the inoculation points. The pathogen was reisolated from

all lesions. Control inoculations with uninoculated MEA disks showed no disease symptoms. Seedlings inoculated with dowels were unaffected 8 mo after inoculation.

Association with bark beetles. The aboveground parts of all trees infected with *V. alacris* were colonized by the European bark beetle *Orthotomicus erosus* (Woll.). *O. erosus* and *Hylurgus ligniperdus* (F.) were present in some of the root crowns, and *Hylastes angustatus* Herbst. and *H. linearis* Erichson were in the roots well below soil level. Isolations from a mixed sample of the two *Hylastes* spp. yielded cultures of *V. alacris*, but the pathogen was not isolated from other bark beetle species.

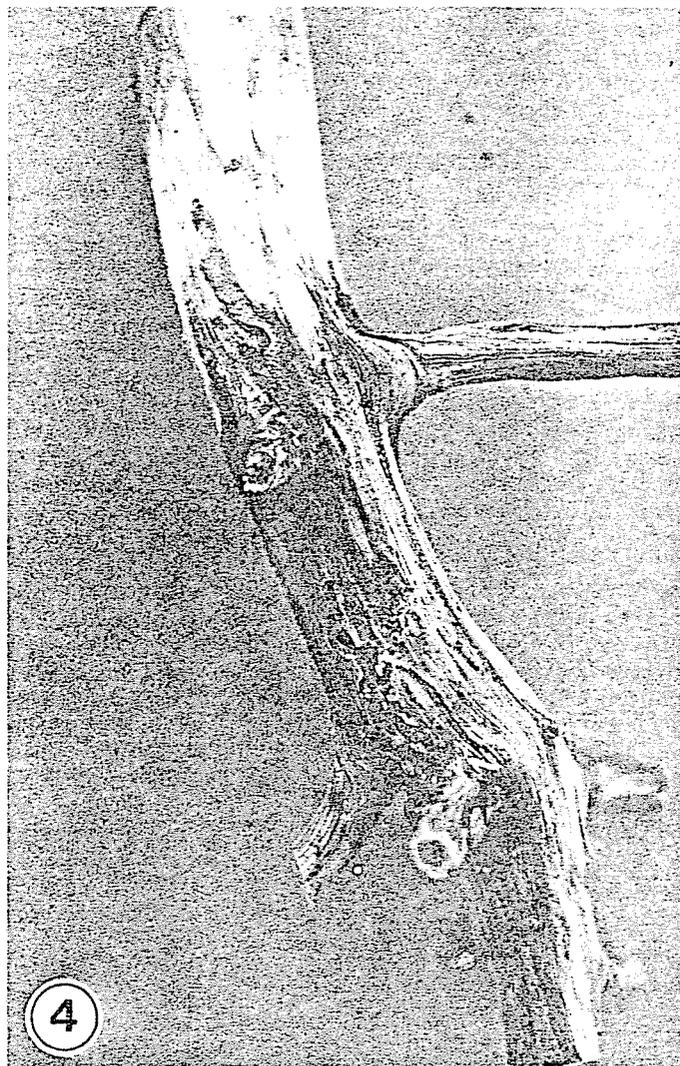
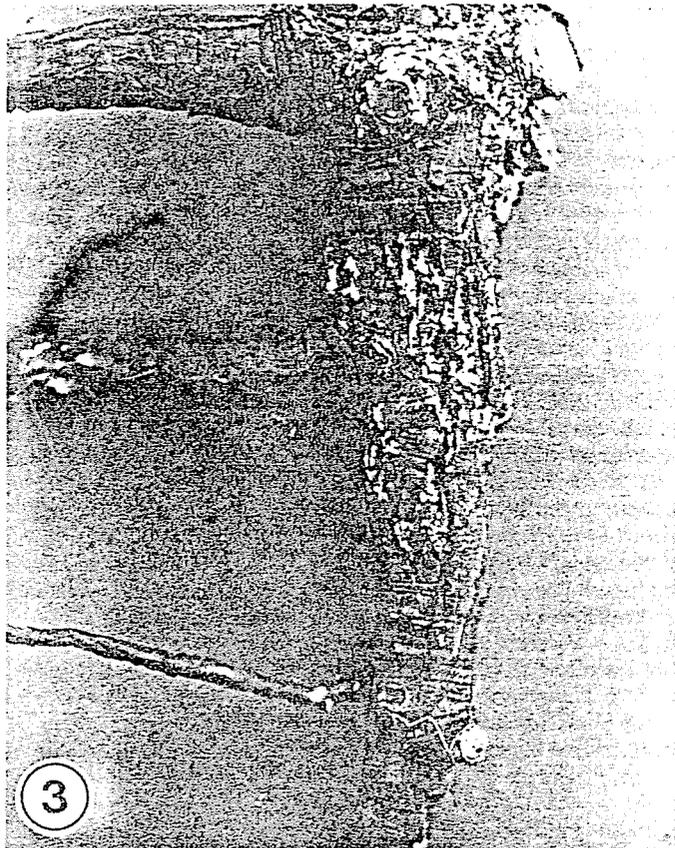
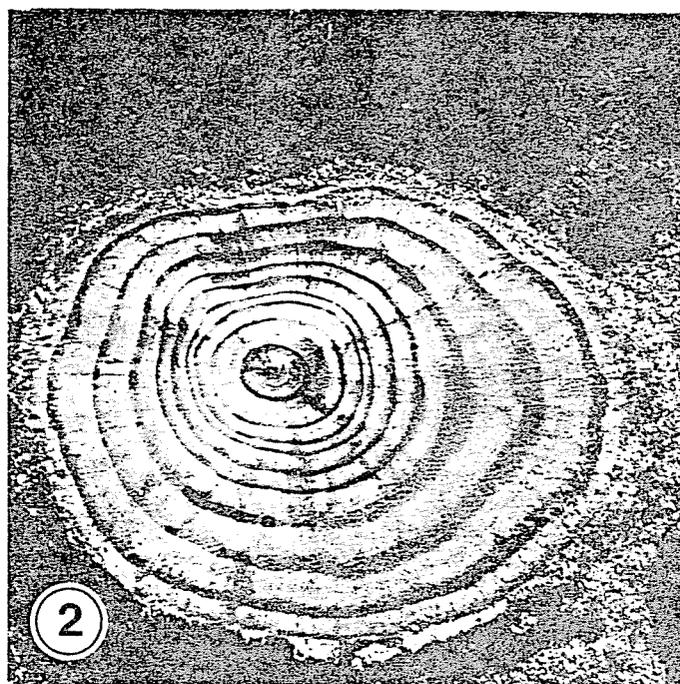
DISCUSSION

Symptoms associated with *V. alacris* were generally similar to those of other *Verticicladiella* diseases (6,15,17,18). However, trees infected with *V. wagnerii* (the best documented *Verticicladiella* root pathogen) usually lose their needles before dying (4), whereas those infected with *V. alacris* retain their needles; and stained sapwood was never found above the root crown area in trees infected with *V. alacris*, whereas trees infected with *V. wagnerii* show extensive discoloration of aboveground parts (4).

V. wagnerii has a low optimum growth temperature in a narrow growth temperature range, and infection is limited by high soil temperatures (16,18). *V. alacris* has a rapid growth rate over a wide temperature range, with an optimum at 25 C (19). The pathogen is favored by the Mediterranean climate of the Western Cape with its cool, wet winters and hot, dry summers.

V. wagnerii infects unwounded roots (2,8), but *V. alacris* and *Verticicladiella* sp. on *P. strobus* L. in New Zealand (15) appear incapable of colonizing uninjured roots. The *V. alacris* infection centers at roadsides suggests that these trees were stressed by disturbance of the soil or damage to the roots (7). Stressed trees (7) and trees with other *Verticicladiella* root diseases (5,9,11) attract bark beetles that carry *Verticicladiella* spp. (3,5,10,14).

Further studies are necessary to establish the relationship between *V. alacris* and bark beetles (particularly *Hylastes* spp.) as possible vectors of this root pathogen and to establish the role of bark beetles such as *O. erosus*, which attack aboveground parts of the trees, in the total disease syndrome.



Figs. 1-4. *Pinus pinaster* and *P. radiata* infected with *Verticillium alacris*: (1) Infection center at a roadside *P. radiata* planting showing excessive production of cones on diseased trees. (2) Staining pattern of wood in the root crown area of an infected *P. pinaster* tree. (3) Resin exudation from the bark of an infected root of a *P. pinaster* sapling. (4) Stained root of an infected *P. pinaster* sapling.

ACKNOWLEDGMENTS

We thank Sharon von Broembsen, Anthea Clarke, D. G. M. Donald, and H. Geertsema for assistance during the investigation. Insect identifications were made by the National Collection of Insects, Pretoria.

LITERATURE CITED

1. BEGA, R. V., D. DOTTA, D. R. MILLER, and R. S. SMITH, Jr. 1966. Root disease survey at Boggs Mountain State Forest, California. *Plant Dis. Rep.* 50:439-440.
2. COBB, F. W., Jr., and W. D. PLATT. 1967. Pathogenicity of *Verticicladiella wagenarii* to Douglas fir. *Phytopathology* 57:998-999.
3. DAVIDSON, R. W., and R. C. ROBINSON-JEFFREY. 1965. New records of *Ceratocystis europhoides* and *C. huntii* with *Verticicladiella* imperfect stages from conifers. *Mycologia* 57:488-490.
4. GOHEEN, D. J. 1976. *Verticicladiella wagenarii* on *Pinus ponderosa*: Epidemiology and interrelationships with insects. Ph.D. thesis, University of California, Berkeley.
5. GOHEEN, D. J., and F. W. COBB, Jr. 1978. Occurrence of *Verticicladiella wagenarii* and its perfect state, *Ceratocystis wagenarii* spp. nov., in insect galleries. *Phytopathology* 68:1192-1195.
6. HALAMBECK, M. 1976. Dieback of Eastern white pine (*Pinus strobus* L.) in cultures. *Polyopr. Znan. Smotra* 39:495-498.
7. HANSEN, E. M. 1978. Incidence of *Verticicladiella wagenarii* and *Phellinus weirii* in Douglas-fir adjacent to and away from roads in western Oregon. *Plant Dis. Rep.* 62:179-181.
8. HELMS, J. A., F. W. COBB, Jr., and H. S. WHITNEY. 1971. Effect of infection by *Verticicladiella wagenarii* on the physiology of *Pinus ponderosa*. *Phytopathology* 61:920-925.
9. HERTERT, H. D., D. L. MILLER, and A. D. PARTRIDGE. 1975. Interaction of bark beetles (Coleoptera: Scolytidae) and root-rot pathogens in grand fir in Northern Idaho. *Can. Entomol.* 107:899-904.
10. KENDRICK, W. B. 1962. The *Leptographium* complex: *Verticicladiella* Hughes. *Can. J. Bot.* 40:771-797.
11. LANE, B. B., and D. J. GOHEEN. 1979. Incidence of root disease in bark beetle-infested eastern Oregon and Washington true firs. *Plant Dis. Rep.* 63:262-266.
12. MILLER, D. L., and A. D. PARTRIDGE. 1974. Root-rot indicators in grand fir. *Plant Dis. Rep.* 58:275-276.
13. PARTRIDGE, A. D., and D. L. MILLER. 1972. Bark beetles and root-rots related in Idaho conifers. *Plant Dis. Rep.* 56:489-500.
14. ROBINSON-JEFFREY, R. C., and A. H. GRIACHENKO. 1964. A new fungus in the genus *Ceratocystis* occurring on blue-stained lodgepole pines attacked by bark beetles. *Can. J. Bot.* 42:527-532.
15. SHAW, C. G., III, and M. DICK. 1980. *Verticicladiella* root disease of *Pinus strobus* in New Zealand. *Plant Dis.* 64:96-98.
16. SMITH, R. S., Jr. 1967. *Verticicladiella* root disease of pines. *Phytopathology* 57:935-938.
17. TOWERS, B. 1976. The occurrence of *Verticicladiella procera* in Pennsylvania. *Plant Dis. Rep.* 61:477.
18. WAGENER, W. W., and J. L. MIELKE. 1961. A staining fungus root disease of ponderosa, jeffrey and pinyon pines. *Plant Dis. Rep.* 45:831-835.
19. WINGFIELD, M. J., and W. F. O. MARASAS. 1980. *Verticicladiella alacris* sp. nov. associated with a root disease of pines in South Africa. *Trans. Br. Mycol. Soc.* In press.

Influence of Cultivar, Age, Soil Texture, and pH on *Meloidogyne incognita* and *Radopholus similis* on Banana

R. G. DAVIDE, Associate Professor of Plant Pathology, College of Agriculture, University of Philippines at Los Baños College, Laguna

ABSTRACT

DAVIDE, R. G. 1980. Influence of cultivar, age, soil texture, and pH on *Meloidogyne incognita* and *Radopholus similis* on banana. *Plant Disease* 64:571-573.

Radopholus similis and *Meloidogyne incognita* were detected in seven banana cultivars, but their population densities differed considerably. Roots of cultivars Giant Cavendish, Cardaba, and Bungulan contained higher populations of *R. similis* than of *M. incognita*; the reverse was true in Dwarf Cavendish, Lacatan, and Latundan. Population densities of both nematode species were low in Saba cultivar. The population of *R. similis* progressively increased and that of *M. incognita* declined with age of Giant Cavendish plantations. This relationship was also observed in plants that had "tip-over" disease and severe root necrosis because of infestation by *R. similis*. Necrotic roots apparently were not suitable for *M. incognita*. Nematodes were present in all soil textures, but both species reproduced better in sandy loam than in soil of finer texture. Population development was most successful at soil pH 5.0-5.6.

Banana (*Musa paradisiaca* L.) is a leading export crop of the Philippines, grown on more than 300,000 ha. The largest banana-producing regions are in the southern and northern Mindanao (6). In a nationwide survey in 1974 and 1975, we found a number of plant-parasitic nematodes associated with different cultivars. The most common species were *Meloidogyne incognita* Chitwood and *Radopholus similis* (Cobb) Thorne, which cause serious damage on banana (1,4). The distribution and population densities of these species varied

considerably in different localities.

This study was done to determine the influence of cultivar, age of plantation, soil texture, and pH on the field distribution and population densities of *R. similis* and *M. incognita*.

MATERIALS AND METHODS

Samples of banana roots and of soil were collected from farms mainly in the provinces of Davao del Norte and Davao del Sur in Mindanao where more than 25,000 ha of Giant Cavendish banana are commercially grown. Approximately 400-cc soil and 4-g (wet weight) root samples per plant hill were randomly collected. All root samples were cut into pieces (1-2 cm long), fixed in FAA for at least 48 hr, and stained for 3-4 min in boiling acid-fuchsin lactophenol.

Stained roots were kept in clear

lactophenol in vials. Later the roots were dissected under a stereomicroscope and the nematodes were identified and counted.

The soil and root samples were collected from the same sites. At each site about 600 cc of soil was obtained approximately 40 cm from the base of the plant at a depth of 15-30 cm. Each soil sample was placed in an individual plastic bag and taken to the nematology laboratory. A 200 cc subsample was sent to the Department of Soil Science for soil identification and pH determination. The results of assays for nematodes other than *R. similis* and *M. incognita* were published elsewhere (3).

RESULTS

Cultivar. The field distribution and population densities of *M. incognita* and *R. similis* varied among cultivars (Fig. 1). The population density of *M. incognita* was greatest in roots of Dwarf Cavendish, Lacatan, and Latundan; the density of *R. similis* was greatest in roots of Giant Cavendish, Bungulan, and Cardaba. Both species were least numerous in Saba.

Age of plantation. The Giant Cavendish plantations in Davao del Norte were from 2 mo to 4 yr old. In plantations less than 1 yr old, the population densities of *M. incognita* and *R. similis* were relatively low and more or less similar (Fig. 2). During the second year, populations increased significantly

This study was supported by a research grant from the National Science Development Board of the Philippines